Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

Claims 1-41 (canceled)

Claim 42 (new): A method for obtaining expression of a tumor suppressor gene 1 2 in a tumor cell in a mammal in vivo, wherein the tumor cell is caused by the absence of a tumor 3 suppressor gene or the presence of a pathologically mutated tumor suppressor gene, the method 4 comprising: 5 contacting the tumor cell with an effective amount of a replication-deficient 6 recombinant adenovirus expression vector comprising: a) a partial or total deletion of a protein 7 IX-encoding DNA sequence, and b) a gene encoding a foreign protein having a tumor suppressive function, wherein said contacting comprises intratumoral, peritumoral or 8 9 intravesicular injection of the recombinant adenenovirus expression vector under suitable 10 conditions such that the foreign protein is expressed in the tumor cell. 1 Claim 43 (new): A method of inhibiting the proliferation of a tumor cell in a 2 mammal, wherein the tumor cell is caused by the absence of a tumor suppressor gene or the 3 presence of a pathologically mutated tumor suppressor gene, the method comprising: 4 administering to the mammal an effective amount of a replication-deficient 5 recombinant adenovirus expression vector comprising: a) a partial or total deletion of a protein 6 IX-encoding DNA sequence; and b) a gene encoding a foreign functional protein having a tumor 7 suppressive function under suitable conditions to the animal, wherein said administering 8 comprises intratumoral, peritumoral or intravesicular injection of the replication-deficient 9 recombinant adenenovirus vector under suitable conditions such that the foreign functional 10 protein is expressed in the tumor cell.

1	Claim 44 (new): The method of claim 42 or 43, wherein the tumor suppressor
2	gene encodes a protein selected from the group p53, p21, p16, Rb, Wilm's tumor WT1 protein,
3	h-NUC, mitosin and mito and p21.
1	Claim 45 (new): The method of claim 42 or 43, wherein the tumor suppressor
2	gene encodes p53.
1	Claim 46 (new): The method of claim 42 or 43, wherein the gene is a suicide
2	gene.
1	Claim 47 (new): The method of claim 42 or 43, wherein the tumor cell is a
2	member selected from the group consisting of non-small cell lung cancer, small cell lung cancer,
3	hepatocarcinoma, melanoma, retinoblastoma, breast tumor, colorectal carcinoma, leukemia,
4	lymphoma, brain tumor, cervical carcinoma, sarcoma, prostate tumor, bladder tumor, tumor of
5	the reticuloendothelial tissues, Wilm's tumor, astrocytoma, glioblastoma, neuroblastoma, ovarian
6	carcinoma, osteosarcoma, or renal cancer.
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1	Claim 48 (new): The method of claim 42 or 43, wherein deletion of the protein
2	IX-encoding DNA sequence extends from about 3500 bp from the 5' viral termini to about 4000
3	bp from the 5' viral termini.
1	Claim 49 (new): The method of claim 42 or 43, wherein the recombinant
2	adenovirus expression vector further comprises a deletion of a non-essential DNA sequence in
3	adenovirus early region 3 or early region 4.
1	Claim 50 (new): The method of claim 42 or 43, wherein the recombinant
2	adenovirus expression vector further comprises a deletion of DNA sequences designated
3	adenovirus E1a and E1b.

Appl. No. 08/958,570 Amdt. dated February 11, 2004 Reply to Office Action of August 11, 2003

l	Claim 51 (new): The method of claim 42 or 43, wherein the recombinant
2	adenovirus expression vector further comprises a deletion of early region 3 or 4 and DNA
3	sequences designated adenovirus E1a and E1b.
İ.	Claim 52 (new): The method of claim 42 or 43, wherein the recombinant
2	adenovirus expression vector further comprises a deletion of up to forty nucleotides positioned 3
3	to the E1a deletion, E1b, protein IX deletions, and wherein said foreign functional protein
1	comprises a polyadenlyation signal.
l	Claim 53 (new): The method of claim 42 or 43, wherein the recombinant
2	adenovirus expression vector is a Group C adenovirus selected from a serotype 1, 2, 5 or 6.
	Claim 54 (new): The method of claim 42 or 43, wherein the recombinant
2	adenovirus expression vector is selected from the group consisting of A/C/N/53 and A/M/N/53.
l	Claim 55 (new): The method claim 42 or 43, further comprising administering a
2	therapeutic agent that controls cell cycle progression and/or induces cell death.
l	Claim 56 (new): The method of claim 42 or 43, wherein the mammal is a
2	human.
l	Claim 57 (new): A method for obtaining expression of a suicide protein in a cell
2	the method comprising administering to the cell an effective amount of a recombinant
3	adenovirus expression vector comprising: a) a partial or total deletion of a protein IX-encoding
1	DNA sequence, and b) a gene encoding a suicide protein, wherein an mRNA encoding the
5	suicide protein is produced by the cell.
ł	Claim 58 (new): A method for reducing the proliferation of a tumor cells in a
2	mammal, the method comprising administering under suitable conditions an effective amount of
3	an adenoviral expression vector comprising: a) a partial or total deletion of a protein IX-

Appl. No. 08/958,570 Amdt. dated February 11, 2004 Reply to Office Action of August 11, 2003

- 4 encoding DNA sequence, and b) a gene encoding a suicide protein or a biologically active
- fragment thereof; and a therapeutic agent that in the presence of the suicide protein is toxic to the
- 6 tumor cell.
- 1 Claim 59 (new): The method of claim 58, wherein the therapeutic agent is a
- 2 thymidine kinase metabolite or a functional equivalent thereof.
- 1 Claim 60 (new): The method of claim 58, wherein the thymidine kinase
- 2 metabolite is ganciclovir or 6-methoxypurine arabinonucleoside or a functional equivalent
- 3 thereof.
- 1 Claim 61 (new): The method of claim 58, wherein the adenoviral expression
- 2 vector is administered by injection into the tumor mass.
- 1 Claim 62 (new): The method of claim 58, wherein the tumor cell is
- 2 hepatocellular carcinoma.
- 1 Claim 63 (new): The method of claim 58, wherein the adenoviral expression
- 2 vector is administered directly into the hepatic artery of the subject.
- 1 Claim 64 (new): The method of claim 58, wherein the cell is present in a
- 2 mammal.
- 1 Claim 65 (new): The method of claim 58, wherein the suicide protein is a
- 2 functional thymidine kinase protein, a functional E. coli DEO \(\Delta \) protein, or a functional cytosine
- 3 deaminase protein.
- 1 Claim 66 (new): The method of claim 58, wherein the recombinant adenovirus
- 2 expression vector further comprises a deletion of a non-essential DNA sequence in adenovirus
- 3 early region 3 or early region 4.

1	Claim 67 (new): The method of claim 58, wherein the recombinant adenovirus
2	expression vector further comprises a deletion of DNA sequences designated adenovirus E1a
3	and E1b.
1	Claim 68 (new): The method of claim 58, wherein the recombinant adenovirus
2	expression vector further comprises a deletion of early region 3 or 4 and DNA sequences
3	designated adenovirus E1a and E1b.
1	Claim 69 (new): The method of claim 58, wherein the recombinant adenovirus
2	expression vector further comprises a deletion of up to forty nucleotides positioned 3' to the E1a
3	deletion, E1b, protein IX deletions, and wherein said foreign functional protein comprises a
4	polyadenlyation signal.
1	Claim 70 (new): The method of claim 58, wherein the recombinant adenovirus
2	expression vector is a Group C adenovirus selected from a serotype 1, 2, 5 or 6.
1	Claim 71 (new): The method of claim 58, wherein the recombinant adenovirus
2	expression vector is selected from the group consisting of A/C/N/53 or A/M/N/53.
1	Claim 72 (new): The method claim 58, further comprising administering a
2	therapeutic agent that controls cell cycle progression and/or induces cell death.
1	Claim 73 (new): The method of claim 58, wherein the tumor cell is a human
2	tumor cell.
1	Claim 74 (new): A kit for reducing the proliferation of tumor cells comprising
2	the components of the adenoviral expression vector of claim 58, a thymidine kinase metabolite
3	or functional equivalent thereof, pharmaceutical carriers and instructions for the treatment of
4	hepatocellular carcinoma using the kit components.